	1. TRANSMITTAL NUMBER:	2. STATE:
TRANSMITTAL AND NOTICE OF APPROVAL OF	0 0 0 2	OKLAHOMA
STATE PLAN MATERIAL FOR: HEALTH CARE FINANCING ADMINISTRATION	3. PROGRAM IDENTIFICATION: TITL SECURITY ACT (MEDICAID)	
TO: REGIONAL ADMINISTRATOR HEALTH CARE FINANCING ADMINISTRATION	4. PROPOSED EFFECTIVE DATE	
DEPARTMENT OF HEALTH AND HUMAN SERVICES	04-08-00	
5. TYPE OF PLAN MATERIAL (Check One):		
□ NEW STATE PLAN □ AMENDMENT TO BE CONS	SIDERED AS NEW PLAN	MENDMENT
COMPLETE BLOCKS 6 THRU 10 IF THIS IS AN AMENE		endment)
6. FEDERAL STATUTE/REGULATION CITATION:	7. FEDERAL BUDGET IMPACT: a. FFY 2000 \$ 8	00.350
1905(a)(12)		90,358 561,437
8. PAGE NUMBER OF THE PLAN SECTION OR ATTACHMENT:	9. PAGE NUMBER OF THE SUPERSE OR ATTACHMENT (If Applicable):	
Attachment 4.19-B, Page 7 Attachment 4.19-B, Page 7a	Same page, Revised 07-01-93 Same page, Revised 10-01-95	
10. SUBJECT OF AMENDMENT:		
Revision and clarification of reimbursement met	chodologies for prescription	druas.
11. GOVERNOR'S REVIEW (Check One): GOVERNOR'S OFFICE REPORTED NO COMMENT COMMENTS OF GOVERNOR'S OFFICE ENCLOSED NO REPLY RECEIVED WITHIN 45 DAYS OF SUBMITTAL	OTHER, AS SPECIFIED:	
12. SIGNATURE OF STATE AGENCY OFFICIAL:	B. RETURN TO:	
13. TYPED NAME:	Oklahoma Health Care Authorit	v
	ttn: Billie Wright	
14. TITLE:	545 N. Lincoln, Suite 124	
Chief Executive Officer	Oklahoma City, OK 73105	
15. DATE SUBMITTED:		
	DE USE ONLY	
17. DATE RECEIVED:	B. DATE APPROVED: September 1	
HIGHT TOUT TO SHEET PLAN APPROVED ON	E COPY ATTACHED	
之间,"我们就是我们,我们就是我们的,我们就是我们的,我们就是我们的,我们就是我们的,我们就是我们的,我们就是我们的,我们就是我们的,我们就是我们的,我们就是	o signature of regional official	يها المالي
21. TYPED NAME: Calvin G. Cline	2 TIME: Associate Regional A Division of Medicald and St	
23. REMARKS:		
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	ou financia de la companie de la co La companie de la co	evi kult. Er kirkt filmlit i a yarkı Arkteriyek verkiyeri illikleri

ICFA

METHODS AND STANDARDS FOR ESTABLISHING PAYMENT RATES OTHER TYPES OF CARE

Payment for prescribed drugs

Payment for compensable drugs is made on the basis of the lowest t the following:

Maximum Allowable Cost (MAC), for state selected products plus a dispensing fee. The State Maximum Allowable Cost (SMAC) is established for certain products which have a Food and Drug Administration (FDA) approved generic equivalent. The products and SMAC price are established based upon the recommendation of the Drug Utilization Review (DUR) Board. The SMAC price is recommended by the DUR to the Agency CEO based on an average of two pricing formulas.

These formulas are as follows: (1) the Oklahoma State and Education Employees Group Insurance Board (OSEEGIB) MAC value and (2) the lower of Average Wholesale Price (AWP) minus 15% or Wholesale Acquisition Cost (WAC) plus 12%. The DUR Board computes formula number one (1) and computes the lower of formula number two (2). The formulas in number one (1) and number two (2) will then be averaged. This will comprise the SMAC price.

(II)Multiple source drugs. Multiple source drug means a drug marketed or sold by two or more manufacturers or labelers or a drug marketed or sold by the same manufacturer or labeler under two or more different proprietary names or both under a proprietary name and without such a name.

Brand Necessary Certification. Unless the prescribing provider certifies a brand name drug product is medically necessary for the well being of the patient, a generic must be substituted for the brand name product. The Brand Necessary Certification applies to the HCFA Upper Limit and State Maximum Allowable Cost (SMAC) products. The requirements for a brand name certification are as follows:

- The certification must be written in the physician's or other prescribing provider's handwriting.
- Certification must be written directly on the prescription blank or on a separate sheet which is attached to the original prescription.
- A standard phrase indicating the need for a specific brand is required. Recommend use of the phrase "Brand Necessary".

Revised 04-08-00

Approval Date 09-19-00 Effective Date 04-08-0

METHODS AND STANDARDS FOR ESTABLISHING PAYMENT RATES OTHER TYPES OF CARE



Payment for prescribed drugs (cont.)

- The printed box on the prescription blank that could be checked by the physician to indicate brand necessary is unacceptable.
- A hand-written statement that is transferred to a rubber stamp and then stamped onto the prescription is unacceptable.

Programming has been developed to review the HCFA upper limit products to assure in the aggregate Medicaid expenditures for multiple source drugs do not exceed the federal upper limits. Such reports and all other relevant statistical data are maintained by the Agency and are available on request.

The Estimated Acquisition Cost (EAC). The EAC means the Agency's best estimate of the price generally and currently paid by providers for a drug marketed or sold by a particular manufacturer or labeler in the package size of drug most frequently purchased by providers. The EAC to be used for the purchase of prescription drug products is established at a percentage of the Average Wholesale Price (AWF) as defined by the Agency's pricing resource. The percentage discount off of the AWP minus is 10.5%.

(IV) The provider's usual and customary charges to the general public. The usual and customary charge will be a single price which includes both the product and the dispensing fee.

After public hearings which considered dispensing fee surveys, usual and customary charge surveys and appropriate inflationary indices, the Agency's Rates and Standards Committee has approved a maximum dispensing fee not to exceed \$4.15.

Claims processed through the MMIS will assure the following:

- Eligibility of patient
- Eligible prescriber
- Eligible participating pharmacist
- Compensability of drug
- Cost within limits
- Limit of prescriptions per month

Revised 04-08-00

Approval Date 991900 Effective Date